## WHAT IS CLAIMED IS:

 A method of detecting colorectal cancer in a human individual comprising: detecting one or more colorectal cancer-associated protein in an extracellular biological sample obtained from a human individual;

wherein the presence of colorectal cancer-associated protein in said extracellular biological sample indicates colorectal cancer in said human individual.

- 2. The method according to Claim 1, wherein said colorectal cancer-associated protein is at least 90% identical to CVA7 or CBF9.
- 3. The method according to Claim 2, wherein said colorectal cancer-association protein is CCA7 or CBF9.
- 4. A method for detecting the presence of a colorectal cancer-associated protein in an extracellular biological sample, the method comprising contacting the biological sample with a binding agent which specifically binds to a colorectal cancer-associated protein selected from the group consisting of CVA7 and CBF9, thereby detecting the presence of the colorectal cancer-associated protein in the extracellular biological sample.
  - 5. The method of Claim 4, wherein the binding agent specifically binds CVA7.
  - 6. The method of Claim 4, wherein the binding agent specifically binds CBF9.
- 7. The method of Claim 4, wherein the biological sample is contacted with a first binding agent that specifically binds CVA7 and a second binding agent that specifically binds CBF9.
- 8. The method of Claim 4, wherein the extracellular biological sample is selected from the group consisting of serum, whole blood, plasma, urine, saliva, sputum, tears, and cerebrospinal fluid.
- 9. The method of Claim 8, wherein the extracellular biological sample is blood or serum.
  - 10. The method of Claim 4, wherein the binding agent is an antibody.
  - 11. The method of Claim 10, wherein the antibody is a monoclonal antibody.
  - 12. The method of Claim 10, wherein the antibody is a polyclonal antibody.
  - 13. The method of Claim 4, wherein the binding agent is bound to a solid support.
  - 14. The method of Claim 13, wherein the solid support comprises nitrocellulose.
- 15. The method of Claim 13, wherein the solid support is a well of a microtiter plate.
  - 16. The method of Claim 4, wherein the binding agent is detectably labled.
  - 17. The method of Claim 16, wherein the label is selected from the group

consisting of a radiolabel, and a fluorescent label.

- 18. The method of Claim 16, wherein the label is a detectable enzyme. 1
- 19. The method of Claim 18, wherein the detectable enzyme is alkaline phosphatase.
- 20. A kit for detecting the presence or absence of a colorectal cancer-associated protein in an extracellular biological sample, the kit comprising a binding agent which specifically binds to a colorectal cancer-associated protein selected from the group consisting of CVA7 and CBF9 and assay reagents for detecting the presence or absence of the colorectal cancer-associated protein in the extracellular biological sample.
  - 21. The kit of Claim 20, wherein the binding agent is labeled.
- 22. The kit of Claim 20, which comprises a first binding agent that specifically binds CVA7 and a second binding agent that specifically binds CBF9.
  - 23. The kit of Claim 20, wherein the binding agent is an antibody.
- 24. The kit of Claim 23, wherein the antibody is a monoclonal antibody or a polyclonal antibody.
  - 25. The kit of Claim 20, wherein the binding agent is bound to a solid support.